<u>Amendments to the Claims</u>: Following is a complete listing of the claims pending in the application, as amended:

- (Currently amended) A process for the preparation of citalopram, comprising characterized in that:
- (a) <u>treating</u> 5-cyanophthalide is treated with a mixture of 4-fluorophenyl magnesium halide and 3-dimethylaminopropyl magnesium halide and, <u>without isolating an intermediate</u>,
- (b) <u>adding the obtained mixture is treated with</u> an organic acid, an inorganic acid, <u>or triphenylphosphine and ethyl azadicarboxylate</u>, a <u>phosphine</u>, or with a labile ester forming group and a base

thereby producing citalogram without isolating an intermediate.

- 2. (Currently amended) A The process according to of claim 1, characterized by the use of using from 1.8 to 2.0 moles of 4-fluorophenyl magnesium halide, for each mole of 5-cyanophthalide.
- 3. (Currently amended) A The process according to of claim 1, characterized by the use of using from 1.09 to 1.2 moles of 3-dimethylaminopropyl magnesium halide, for each mole of 5-cyanophthalide.
- 4. (Currently amended) A The process according to of claim 1, characterized by the use of using from 1.7 to 1.6 moles of 4-fluorophenyl magnesium halide, are used for each mole of 3-dimethylaminopropyl magnesium halide.

- 5. (Currently amended) A <u>The</u> process according to <u>of</u> claim 1, characterized by the fact that wherein the 4-fluorophenyl magnesium halide is a bromide.
- 6. (Currently amended) A <u>The</u> process according to <u>of</u> claim 1, characterized by the fact that <u>wherein the</u> 3-dimethylaminopropyl magnesium halide is a chloride.
- 7. (Currently amended) A <u>The</u> process according to of claim 1, characterized by the fact that said wherein the acid has a pK comprised from 0 to 3.
- 8. (Currently amended) A <u>The</u> process according to <u>of</u> claim 1, characterized by the fact that said <u>wherein the</u> acid has a pK comprised from 2 to 3.
- 9. (Currently amended) A <u>The</u> process according to <u>of</u> claim 7, characterized by the fact that said <u>wherein the</u> acid is ortoortho-phosphoric acid.
- 10. (Currently amended) A <u>The</u> process according to <u>of</u> claim 7, characterized by the fact that <u>wherein</u> the acid is used in a concentration comprised from 55 to 95% by weight, preferably in concentration of about 85% by weight.

11-15. (Canceled)

16. (Currently amended) A <u>The process according to of claim 1, characterized by the fact that the process is carried out in an organic polar aprotic solvent.</u>

- 17. (Currently amended) A <u>The</u> process according to <u>of</u> claim 16, characterized by the fact that the process is carried out in from 1.0 to 1.6 litres of solvent, for each mole of 5-cyanophthalide.
- 18. (Currently amended) A <u>The</u> process according to <u>of</u> claim 16, characterized by the fact that wherein the solvent is selected from tetrahydrofuran and/or toluene.
- 19. (Currently amended) (Currently amended) A <u>The</u> process according to <u>of</u> claim 1, characterized by the fact that the step (a) is carried out at 20/+20 -20 to +20° C.
- 20. (Currently amended) A <u>The</u> process according to <u>of</u> claim 1, characterized by the fact that the wherein step (a) is carried out at -10/0 -10 to 0° C.
- 21. (Currently amended) A <u>The</u> process according to <u>of</u> claim 1, characterized by the fact that the wherein step (b) is carried out at -10/+20 -10 to +20° C.
- 22. (Currently amended) A <u>The</u> process according to <u>of</u> claim 1, characterized by the fact that the wherein step (b) is carried out at 0/+10 0 to +10° C.
 - 23. (Canceled)
 - 24. (Currently amended) A Compound of formula:

where X is an <u>a</u> halogen, preferably chlorine or bromine.

- 25. (Canceled)
- 26. (New) The compound of claim 24, wherein X is chlorine or bromine.
- 27. (New) A one pot process for the preparation of citalopram, comprising:
 combining 5-cyanophthalide 4-fluorophenyl magnesium halide and 3dimethylaminopropyl magnesium halide in a pot, and, without isolating an intermediate,
 performing acid catalysed cyclization, thereby producing citalopram in one pot without
 isolating an intermediate.